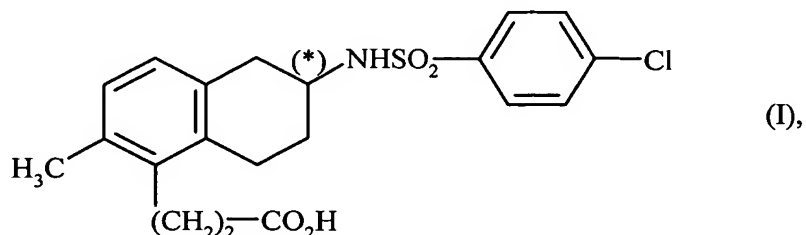


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LISTING OF CLAIMS

Claims 1-12 (CANCELED)

13- (NEW) A solid orodispersible pharmaceutical composition comprising compound
 5 A of formula (I), optionally in the form of an optical isomer, or a pharmaceutically acceptable salt thereof :



and granules consisting of co-dried lactose and starch.

10 14- (NEW) The composition according to Claim 13, wherein the composition disintegrates in the mouth in less than three minutes.

15- (NEW) The composition according to Claim 14, wherein the composition disintegrates in the mouth in less than one minute.

16- (NEW) The composition according to Claim 13, wherein compound A has the (R) configuration.

15 17- (NEW) The composition according to Claim 13, comprising, in relation to the total weight of the composition :

- from 2.5 % to 20 % by weight of compound A or a pharmaceutically acceptable salt thereof, and
- from 75 % to 95 % by weight of granules consisting of co-dried lactose and starch.

20 18- (NEW) The composition according to Claim 17, comprising from 5 % to 10 % by weight of compound A or a pharmaceutically acceptable salt thereof.

19- (NEW) The composition according to Claim 13, wherein compound A is in the form of a sodium salt.

20- (NEW) The composition according to Claim 13, further comprising one or more flavourings and sweeteners.

5 **21- (NEW)** The composition according to Claim 13, further comprising one or more lubricants and a flow agent.

22- (NEW) The composition according to Claim 13, wherein the composition is in the form of a tablet.

10 **23- (NEW)** The tablet according to Claim 22, wherein the tablet is obtained by direct compression.

24- (NEW) The tablet according to Claim 23, wherein the tablet has a hardness from 15 to 30 Newtons.

15 **25- (NEW)** A process for the manufacture of solid orodispersible compositions of compound A, or a pharmaceutically acceptable salt thereof, which disintegrate in the mouth in less than three minutes, wherein compound A, or a pharmaceutically acceptable salt thereof, is mixed with granules consisting of co-dried lactose and starch.

20 **26- (NEW)** A process for the manufacture of solid orodispersible compositions of compound A, or a pharmaceutically acceptable salt thereof, which disintegrate in the mouth in less than one minute, wherein compound A, or a pharmaceutically acceptable salt thereof, is mixed with granules consisting of co-dried lactose and starch.

27- (NEW) A method for treating a living animal body, including a human, afflicted with a condition treatable by an antithrombotic agent, comprising the step of administering to the living animal body, including a human, a composition according to claim 13 which is effective for alleviation of the condition.